CLEAN VERSION OF REWRITTEN OR ADDED CLAIMS
PURSUANT TO 37 C.F.R. §1.121 (c)(1)(i)

Please add the following claims:

34. A method of treatment, comprising:
    a) providing:
        i) a mammal having symptoms of sepsis,
        ii) a therapeutic preparation, consisting of anti-TNF-α and anti-IL-6 antibodies, and one or more inactive ingredients; and
        iii) administering said preparation to said mammal wherein said symptoms are reduced.

35. The method of Claim 34, wherein said inactive ingredient is bovine serum albumin.

36. The method of Claim 34, wherein said mammal is a human.

37. The method of Claim 34, wherein said administering is performed intravenously.

38. The method of Claim 34, wherein said administering is performed orally.

39. The method of Claim 34, wherein said administering is performed parenterally.

40. The method of Claim 34, wherein said antibodies are polyclonal antibodies.

41. The method of Claim 40, wherein said polyclonal antibodies are avian antibodies.

42. A method of treatment, comprising:

a) providing:

i) a mammal having symptoms of sepsis,

ii) a therapeutic preparation, comprising polyclonal anti-TNF-α and polyclonal anti-IL-6 antibodies; and

iii) administering said preparation to said mammal wherein said symptoms are reduced.

F1 cont.

43. The method of Claim 42, wherein said therapeutic preparation further comprises anti-IFN antibodies.

44. The method of Claim 42, wherein said mammal is a human.

45. The method of Claim 42, wherein said administering is performed intravenously.

46. The method of Claim 42, wherein said administering is performed orally.

47. The method of Claim 42, wherein said administering is performed parenterally.

48. The method of Claim 42, wherein said polyclonal antibodies are avian antibodies.